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Silencing of Germline-Expressed Genes by DNA Elimination in Somatic Cells

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SUMMARY

Chromatin diminution is the programmed elimination of specific DNA sequences during development. It occurs in diverse species, but the function(s) of diminution and the specificity of sequence loss remain largely unknown. Diminution in the nematode *Ascaris suum* occurs during early embryonic cleavages and leads to the loss of germline genome sequences and the formation of a distinct genome in somatic cells. We found that ~43 Mb (~13%) of genome sequence is eliminated in *A. suum* somatic cells, including ~12.7 Mb of unique sequence. The eliminated sequences and location of the DNA breaks are the same in all somatic lineages from a single individual, and between different individuals. At least 685 genes are eliminated. These genes are preferentially expressed in the germline and during early embryogenesis. We propose that diminution is a mechanism of germline gene regulation that specifically removes a large number of genes involved in gametogenesis and early embryogenesis.

INTRODUCTION

Metazoans must both ensure the stability of their genomes, and also carefully regulate the expression of germline genes in somatic tissues. Failure of either of these processes has

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SUPPLEMENTAL INFORMATION

Supplemental Information includes 4 figures, 6 tables, and Supplemental Experimental Procedures.

Accession numbers and URLs

Genomic and RNA-seq reads were deposited at the SRA (SRP013430, SRP013609 and ERP000532), GSS (DU976516.1 - DU978593.1, ED033836.1 - ED479302.1, ED562826.1 - ED567026.1), and GEO (GSE38470) databases. The germline and somatic genomes are also available in NCBI BioProject database with accession number PRJNA62057. *Ascaris suum* genome browser: <http://ascaris.nematodegenomes.org/>. Data is also deposited at <http://nematode.net> and <http://www.wormbase.org/>.

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severe consequences. However, there are examples of programmed genome instability that are integral to the biology of an organism. Well-known examples include vertebrate immunoglobulin gene rearrangement that enables antibody diversification (Jung et al., 2006) and the extensive remodeling of the somatic genome that occurs during development of the macronucleus in ciliates (Chalker and Yao, 2011). Chromatin diminution is another form of genome rearrangement with DNA loss that occurs during the development of diverse Metazoa including some nematodes, copepod crustaceans, insects, lampreys/hagfish, and zebra finches (Bachmann-Waldmann et al., 2004; Goday and Esteban, 2001; Goday and Pimpinelli, 1993; Kloc and Zagrodzinska, 2001; Muller et al., 1996; Muller and Tobler, 2000; Nemetschke et al., 2010; Smith et al., 2009; Tobler et al., 1992; Tobler et al., 1985; Zufall et al., 2005). DNA elimination occurs either during gametogenesis or during differentiation of the somatic lineage early in embryonic development. In some species, DNA elimination may be involved in sex determination (Goday and Esteban, 2001; Goday and Pigozzi, 2010; Nemetschke et al., 2010), but in most organisms that undergo programmed chromatin diminution, the sequences lost and the role(s) of diminution remain unknown.

Chromatin diminution was first described in an ascaridid nematode by Theodor Boveri in 1887 (Boveri, 1887). Diminution in nematodes is restricted to a number of parasitic nematodes primarily in the Ascarididae (Goday and Pimpinelli, 1993; Muller et al., 1996; Muller and Tobler, 2000; Pimpinelli and Goday, 1989) and does not occur in *Caenorhabditis elegans* (Emmons et al., 1979). In the pig parasite *A. suum*, diminution occurs during the third through fifth cleavages (4 to 16 cell stage) of development in five distinct somatic precursor cells that give rise to different cell lineages (Fig. 1). This raises the key questions: 1) Are the sequences lost and the changes that occur during diminution the same in all five of these precursor cells, and 2) What are the sequences that are lost and their functional significance?

During the *Ascaris* diminution process, chromosomes are broken and the fragments to be eliminated remain at the metaphase plate while the retained DNA is segregated into daughter cells. It has been estimated that 25% of *A. suum* genome is lost in somatic cells, whereas the germline genome remains intact (Muller et al., 1996; Muller and Tobler, 2000; Tobler et al., 1992; Tobler et al., 1985). The eliminated DNA includes highly repetitive satellite sequences consisting primarily of a 121 bp tandem repeat that is located in heterochromatin-like blocks at internal sites and chromosome ends (Muller et al., 1982; Niedermaier and Moritz, 2000; Streeck et al., 1982). Both internal regions and terminal heterochromatic regions are eliminated, with addition of new telomeres resulting in a ~50% increase in chromosome number (Bachmann-Waldmann et al., 2004; Huang et al., 1996; Jentsch et al., 2002; Magnenat et al., 1999; Muller et al., 1991; Niedermaier and Moritz, 2000). While a few chromosomal breakpoints where new telomere addition occurs have been partially characterized (Bachmann-Waldmann et al., 2004; Huang et al., 1996; Jentsch et al., 2002; Magnenat et al., 1999; Muller et al., 1991) and three single-copy genes that are eliminated have been identified (Etter et al., 1994; Huang et al., 1996; Spicher et al., 1994), the full complement of eliminated genes and sequences and the consequences for the somatic cells remain unknown (Goday and Pimpinelli, 1993; Muller et al., 1996; Muller and Tobler, 2000; Pimpinelli and Goday, 1989).

RESULTS and DISCUSSION

Identification of Eliminated Sequences and Breakpoints Associated with *Ascaris* Diminution

We deep sequenced genomic DNA libraries from the spermatids (germline) and intestine (somatic) of a single male *A. suum* (Table S1) to enable a genome-wide analysis of DNA

changes following diminution. We used these data to independently assemble the germline and somatic genomes (Table 1), and mapped the raw reads of the germline and somatic genomes back to the germline assembly. These analyses identified sequences that are eliminated to form the somatic genome and DNA breakpoints associated with *A. suum* chromatin diminution (Fig. 2 and Fig. S1). Paired-end and mate-pair reads for the germline and somatic regions strongly supported the germline assembly, the identified breakpoints, and DNA loss (Tables S3 and S4). The results indicated that the major type of genome alteration is chromosome breakage, loss of DNA sequence, and the healing of retained chromosomes by telomere addition (Figs. 3 and 4 and Tables S3 and S4). We found no evidence for the loss of interstitial sequences followed by DNA fusions or other genome rearrangements. Breakpoints, DNA loss, and new telomere addition identified in the genome sequencing data were confirmed using specific PCR assays (Figs. 2 and 3). Among the sequences eliminated are at least 35 loci that may have arisen through recent duplication and rearrangement in the *A. suum* lineage (Fig. 4).

Diminution Process is Conserved in Distinct Somatic Lineages and Between Male and Female

To determine if diminution happens in the same way in all somatic cell lineages (Fig. 1), we compared the sequences lost from the male intestine to those lost from other somatic cell types (e.g., body wall muscle, pharyngeal muscle, hypodermis, and neurons) (Table S1) in the same male. The intestine is derived from a single cell (E) in the *A. suum* cell lineage (Fig. 1) whereas other somatic tissues are derived from additional and often multiple cell lineages that independently undergo chromatin diminution (Fig. 1, cells labeled AB, C, and D) (Boveri, 1899, 1910). We found that the DNA loss and chromosome breaks in an individual male are conserved between the intestine and other somatic cell types (Fig. 3D and Fig. S1), suggesting that the mechanism and consequences of diminution are the same in different cells. Additional comparison of the sequence lost between the male and a female worm shows a high degree of fidelity in the breaks and DNA loss between individuals (Fig. 3D and Fig. S1). However, there is some heterogeneity in the exact position, with ~80% of the breaks occurring within 500 bp of each other within an individual and ~70% within 1,000 bp between individuals (Fig. 3E and Fig. S1) consistent with earlier studies (Bachmann-Waldmann et al., 2004; Huang et al., 1996; Jentsch et al., 2002; Magnenat et al., 1999; Muller et al., 1991). Analysis of DNA sequence up to 5 kb on either side of the telomere addition sites did not reveal any specific sequence motifs or other characteristics that might mark the regions for chromosomal breakage (Fig. S2 and Supplementary Experimental Procedures).

A Large Number of Germline Genes are Eliminated During Diminution

Our analysis of the DNA lost revealed that ~43 Mb (~13%) of sequence is eliminated from the germline genome during the formation of the somatic genome (Table 1). The majority of the eliminated sequence (29 Mb, 70% of the eliminated sequence) is the 121 bp satellite, repeat sequence previously described (Muller and Tobler, 2000; Muller et al., 1982; Streeck et al., 1982) (Table S2). No other major loss of repetitive sequence was observed. The remaining ~12.7 Mb of the eliminated sequence is unique and includes at least 685 predicted genes (Tables S5 and S6). We sought to identify whether the eliminated genes shared any common features in their patterns of expression. Genome-wide expression profiles were constructed by RNA-seq of poly(A)⁺ RNA from testis, ovary, embryo, larvae, intestine, muscle and other somatic tissues (Table S1), and the expression level was measured by reads per kilobase of template per million mapped reads (RPKM). Remarkably, these results revealed that 85% of the eliminated genes are expressed preferentially during gametogenesis or early embryogenesis, and the remaining 15% are expressed in both (Fig. 5 and Table S6). Most of these genes are expressed specifically in the testis and therefore are likely involved

in spermatogenesis. However, not all genes expressed in the germline were eliminated during chromatin diminution. Functional annotations of the eliminated genes suggest that they are enriched for protein kinases, protein phosphatases, proteins associated with chromatin, RNA and nucleotide binding proteins, and translation initiation proteins (Table S6). These proteins are associated with network functions including protein synthesis, RNA post-transcriptional modification, gene expression, cell death, and cellular compromise (Table S6). Notably, at least 30 families of these eliminated genes are orthologs of well-characterized genes in *C. elegans* whose loss is associated with clear phenotypes in germline formation, gametogenesis, and early embryogenesis (e.g., *air-1*, *gld-2*, *cgh-1*, *gla-3*, *fer-1*, *spe* genes, and *pab* genes) (Table S6).

Eliminated Genes Suggest Biological Functions for Diminution

We observed that ~53% (363) of the eliminated genes have paralogs in the genome (Table S6). This is consistent with a model where an ancient genome duplication in the *A. suum* lineage was balanced by chromatin diminution to regulate gene dosage or to provide a mechanism for the selective retention of specific genes and thus their function (Bachmann-Waldmann et al., 2004; Goday and Esteban, 2001; Goday and Pimpinelli, 1993; Muller et al., 1996; Muller and Tobler, 2000; Tobler et al., 1992; Tobler et al., 1985). A previous study demonstrated one of the two paralogs of ribosomal protein rps-19 gene was eliminated in *A. suum*, suggesting that the two proteins may play differential roles in translation (Etter et al., 1994). Recent data on rps-19 indicates that mutations or knockdown of this and other specific ribosomal protein genes in vertebrates leads to discrete changes in the translation of specific mRNAs, but not general translation (Horos et al., 2012; Kondrashov et al., 2011). Our observation that a major group of the eliminated genes is associated with translation (Tables S6) reinforces the idea that the translation machinery may differ between the germline and soma. Notably, we found that a number of translation initiation factors are eliminated from the germline (eIF4E, eIF5, eIF2 subunit 2, eIF3B, eIF3C, eIF3D, eIF3i, and eIF2 subunit 3). Another, non-exclusive model of the function of diminution is that in addition to the elimination of the genes, the chromosomal regions eliminated play an important role in chromatin organization that contribute to broader gene regulation. For example, these regions may repress genes in the germline or their elimination may activate key somatic genes. We found no evidence of telomeric position effect silencing of genes due to telomere addition (Huang et al., 1996) (Fig. S1) but cannot eliminate the possibility of other indirect effects.

Ascaris Chromatin Diminution, Small RNAs, and Marks for Chromosome Breaks

In ciliates, small RNAs (piRNAs) are known to play a key role in the programmed DNA rearrangements and elimination (Chalker and Yao, 2011). We looked previously for small RNAs (Wang et al., 2011) that might be associated with *Ascaris* chromatin diminution, in particular RNAs mapping to the 121-bp repeat element that constitutes ~30 Mb of the eliminated sequence. We characterized small RNAs before, during, and after chromatin diminution and found no correlation between the eliminated repeats and any small RNAs. Furthermore, piRNAs and PIWI Argonautes are absent in *Ascaris* (Wang et al., 2011). Recent studies in *C. elegans* suggest that small RNAs also play a role in the recognition of “self” versus “non-self” and in multigenerational epigenetic inheritance (Ashe et al., 2012; Bagijn et al., 2012; Buckley et al., 2012; Lee et al., 2012; Luteijn et al., 2012; Shirayama et al., 2012). We reexamined the expression of *Ascaris* small RNAs (Wang et al., 2011) correlating with chromatin diminution. We found no temporal or any other correlation of small RNAs associated with regions retained, regions eliminated, or the chromosome breakpoint regions. While preliminary analyses of *Ascaris* polyA+ RNA levels (RNA-seq) demonstrate that several Argonaute and other protein mRNAs increase during the period of early embryo development leading up to and during the time of diminution, their increased

expression may not be correlated with diminution and could be associated with the maternal to zygotic transition or serve other functions during this complex period of development. Overall, these and other data in *Ascaris* [including the loss of large numbers of genes, the lack of discrete sequence elements that mark sites of DNA breaks for telomere addition (Fig. S2), and the absence of removal of interstitial DNA sequences followed by DNA fusion] suggest that the function and mechanism for DNA elimination in *Ascaris* may differ from the programmed rearrangements in ciliates. As we did not identify discrete sequence elements that mark the sites of DNA breaks, we suggest that epigenetic marks (and even small RNAs yet to be identified) could play an important role in defining chromosome break sites and play a role in chromatin diminution. Additional studies will be required to further examine these possibilities.

Recently, a preliminary genomic study on the programmed DNA elimination in sea lampreys also demonstrated that unique sequences were eliminated from somatic cells (Smith et al., 2012). Among the genes eliminated in the somatic cells were some involved in transcriptional programs that are likely to play a role in maintaining germline function. Thus, elimination of specific germline-expressed genes in metazoa may be a common function of chromatin diminution.

Our work is a comprehensive analysis of the germline and somatic genome from a metazoan, the DNA lost and the chromosome changes that occur, and the elimination of specific germline expressed genes suggesting a function for that *Ascaris* chromatin diminution and a paradigm for DNA elimination. A hallmark of most Metazoa is that germline cells are set aside early in development. Soma-specific elimination provides a unique mechanism of gene repression, reminiscent of Weismann's original theory of the differentiation between germline and soma (Weissmann, 1893). Our comprehensive identification of the genome changes in the soma of *A. suum* provides the foundation for the elucidation of the features and epigenetic changes underlying the mechanisms of selective DNA breakage and DNA loss in chromatin diminution. Understanding chromosome breaks, telomere healing, and selective DNA loss in chromatin diminution is likely to offer insight into genome stability and changes in normal processes and disease.

EXPERIMENTAL PROCEDURES

Sample collection and DNA isolation

A. suum samples were collected from pig intestines at a slaughterhouse in Sandusky, OH, U.S or Ghent, Belgium. A single male *A. suum* (U.S.) was dissected and the spermatids, the intestine, the testis, and the remaining tissue (carcass, which includes muscle, hypodermis, pharynx, and neurons) collected, frozen in liquid nitrogen, and stored at -80°C . A single female (U.S.) was dissected and the ovary/oviduct, uterus, intestine, and carcass collected, frozen in liquid nitrogen, and stored at -80°C .

Genome sequencing and assembly

Sequencing—Genomic DNA libraries were constructed from *A. suum* germline and somatic tissues and sequenced (Table S1). DNA isolation and libraries were constructed using standard methods and Illumina protocols and sequenced on the Illumina GAIIx or HiSeq platforms except where noted in the supplemental Materials and Methods. Genomic reads for assembly, scaffolding and analysis are listed in Table S1. The fold coverage numbers in Table S1 for all these libraries are derived from high quality reads that can be mapped back to the final assemblies.

Assembly—Reads from germline or somatic sources were used to independently assemble the two genomes. To minimize the sequence heterogeneity, we only used DNA sequences from a single male for the generation of consensus sequences within the assembly. Genomic reads for scaffolding (Table S1) were only used to confirm and support the links that bridge contigs into scaffolds, and none of these sequences were incorporated into the genome assemblies. Because of the presence of some duplicated loci (Fig. 4 and Table S4), we applied a “sub-assembly” strategy to capture all changes that occur in the germline genome. First, we built an initial germline genome assembly using velvet (v1.1.03) (Zerbino and Birney, 2008). From this assembly, we defined ~12.7 Mb of eliminated sequences (see below). All mappable germline reads were divided into 2 groups: those reads to the 12.7 Mb eliminated regions and all their pairs (from the paired-end and mate-pair libraries) and those reads retained in the soma following diminution. Next, we assembled the two groups of reads independently by using velvet (Zerbino and Birney, 2008). Finally, we combined the eliminated and retained assemblies using phrap (<http://www.phrap.org/>, v1.080812). Each assembly step was optimized and scaffolding performed under the overall guideline of sequentially bridging gaps 200, 500, 1,500, and 3,500 bp with 20 pairs of supporting reads, and the assemblies were checked using the Tablet assembly viewer (v1.11.11.01). The somatic genome was built using velvet, with 10 pairs of paired-end Illumina reads or 3 pairs of Sanger capillary reads for scaffolding, and the assembly also checked using the Tablet viewer. Note the N50 of the somatic assembly is not as good as that of the germline. This is mainly due to the lower average sequencing depth on the large fragment paired-end library for the somatic genome (compared with the germline) available for scaffolding.

Identification of eliminated sequences, germline genome changes, and breakpoints with telomere addition

We used two libraries generated and analyzed in parallel with similar coverage (2×150 bp, 360 bp insert size, $\sim 50 \times$ coverage) from the single male spermatids and intestine to map reads back to the germline genome assembly (bowtie2, v2.0.0-beta5) (Langmead and Salzberg, 2012) to identify eliminated sequence and genome changes in the germline. We used a 100-bp window to scan the genome for regions with a somatic/germline coverage ratio < 0.1 to provide an initial set of long sequence blocks containing potentially eliminated sequences. To identify germline genome changes, we manually checked all scaffolds 3 kb with 500 bp contiguous coverage. We identified 102 loci with DNA alterations (Tables S3 and S4). We then compared their sequences (10 Kb flanking the changes) to the somatic genome assembly, identified their exact positions where DNA loss occurred, and used these positions to establish the eliminated and retained regions in scaffolds. Germline scaffolds that did not harbor any DNA breakpoints were defined either as retained or eliminated based on the coverage ratio (see Fig. 2A and Fig. S1 for examples). From the somatic assembly, we also identified DNA breakpoints with addition of telomeres. Somatic scaffolds with telomeric sequences were independently confirmed by paired-end reads (Table S3) and PCR (Fig. 3A–C). For those DNA changes without telomere addition, their somatic loci were evaluated by germline paired-end reads (Table S4) and PCR analysis (Fig. 4) to confirm their presence in the germline.

RNA preparation, RNA-seq and assembly

Samples for RNA preparations are the same as those described for DNA preparations above or previously (Wang et al., 2011). Total RNA was prepared and RNA-seq libraries were made and sequenced as described (Wang et al., 2011). For each sample, 200 μ g of total RNA was used for poly(A) selection and 200 ng of poly(A)+ RNA was used to make the cDNA libraries. Two cDNA assemblies were made by using all the RNA-seq data: one is a de-novo assembly using velvet/oases (v1.1.03/v0.2) (Zerbino and Birney, 2008), which has been used for the genome functional coverage assessment and gene annotation pipeline; the

other is map-based assembly using tophat/cufflinks (v1.3.0) (Trapnell et al., 2012), which was used to facilitate gene prediction (see below).

Gene model prediction

We integrated gene evidence from multiple sources to build gene models for the *A. suum* germline genome. First, a two-pass MAKER (Holt and Yandell, 2011) annotation pipeline (v2.22) was used. In the 1st pass, evidence was used from the RNA-seq assembly, alignments to the Swiss-prot protein database, predictions of the *ab initio* gene finders SNAP (v2010-07-28) (Korf, 2004), and trained using CEGMA (v2.0) (Parra et al., 2007) gff output and GeneMark-ES (v2.3e) (Lomsadze et al., 2005). For the 2nd pass, first-pass MAKER gff files were used to train Augustus (v2.5.5) (Stanke and Waack, 2003), retrain the SNAP models, and MAKER was rerun with the addition of these two programs. Second, we annotated genomic regions without MAKER genes by RNA-seq data using tophat/cufflinks (v1.3.0) (Trapnell et al., 2012). Last, regions without MAKER and tophat/cufflinks genes were further annotated with transferred annotations from a published *A. suum* assembly (Jex et al., 2011) by using RATT (Otto et al., 2011). The final gene set consists of 11,446 genes from MAKER, 2,947 genes from tophat/cufflink and 1,053 genes from RATT.

Gene expression analysis

To profile the tissue expression of all *A. suum* genes, we used 8 different developmental stages/tissues of *A. suum*, including testis, ovary, embryo, larvae, intestine, muscle, male carcass and female carcass (Table S1). For each predicted gene, their expression level (RPKM) was calculated using tophat/cufflinks (Trapnell et al., 2012). Genes with a RPKM ≥ 2 and the RPKM ≥ 1.5 fold higher in one particular tissue compared to all other tissues were defined as enriched gene expression in a particular tissue (maternal genes RPKM $\geq 2/3 \times$ higher than the embryo) (Figs. 5 and Table S5). This is a relatively high stringency cutoff due to the existence of neighbor/similar tissues in development, such as ovary/embryo/larvae and muscle/male carcass/female carcass in this analysis. The expression profiles for tissue specific genes and other genes were clustered by using Cluster (v3.0) (Eisen et al., 1998) and visualized in heatmaps using treeview (<http://jtreeview.sourceforge.net/>) (Fig. 5). For these analyses, the average expression in muscle, male carcass, and female carcass was used to estimate the baseline expression level for other somatic tissues. Groups of enriched genes are also illustrated in dotplots for enriched tissue versus all other tissues (Fig. S3).

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Highlights

- Over 685 (5%) of *Ascaris suum* genes are eliminated during chromatin diminution
- Eliminated genes are mainly expressed during gametogenesis and early embryogenesis
- The genome changes are conserved in different somatic cell lineages and sexes
- Diminution may be a mechanism of gene regulation that silences germline genes

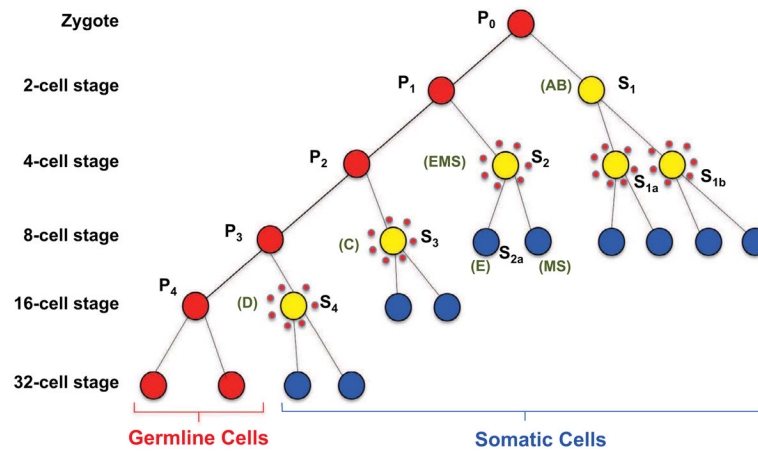


Figure 1. *A. suum* early embryo development, cell lineage and chromatin diminution

Primordial germ cells (P) are in red, cells undergoing chromatin diminution are represented by yellow filled circles surrounded by dots, and blue cells (S) are precursor somatic cells and lineages. The primordial germ cells numbers correspond to their division state. P₀ is the zygote whereas P₁ through P₄ represent the primordial germ cell derived from each subsequent cleavage of the germ cells as illustrated. S₁–S₄ cells are successive precursor somatic cell derived from each division of a germ cell (EMS = intestine, body wall muscle, and pharynx; E = intestine; MS = Body wall muscle, neurons, somatic gonad, coelomocytes, and pharynx; AB = Nervous system, hypodermis, and pharynx; C = Body wall muscle, hypodermis, and neurons; and D = Body wall muscle). Adapted from Theodor Boveri (Boveri, 1899, 1910) and Fritz Müller and Heinz Tobler (Goday and Pimpinelli, 1993; Muller et al., 1996; Muller and Tobler, 2000; Pimpinelli and Goday, 1989).

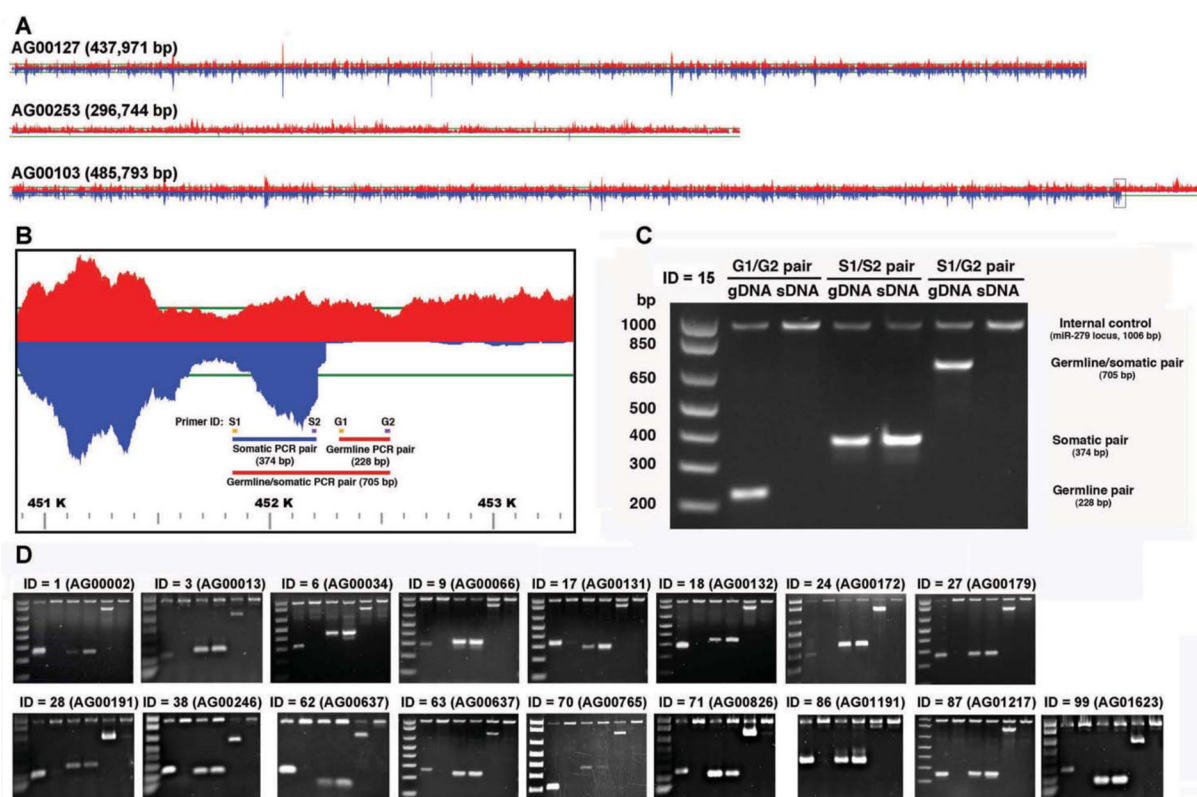


Figure 2. *A. suum* DNA elimination

A. Germline and somatic read coverage for regions of the *A. suum* genome illustrating the retention of segments in both the germline and somatic tissue (top), a region completely eliminated in the somatic tissue (middle), and a DNA breakpoint and region eliminated in the somatic tissues (bottom). Red designates germline reads and blue are somatic reads with the horizontal green line representing 50-fold coverage.

B. Enlarged region of a scaffold (AG00103, Fig. 2A bottom) illustrating the PCR strategy used to verify DNA elimination predicted from the comparison of the germline and somatic genome sequences.

C. PCR data confirm the elimination of DNA corresponding to scaffold AG00103 in *A. suum* somatic tissues. Note that the germline primer pair (G1/G2) produces a PCR product in the germline DNA (gDNA), but not somatic DNA (sDNA). The somatic primer pair (S1/S2) leads to PCR products in both the germline and somatic DNA, and the primer pair spanning the breakpoint (S1/G2) produces a PCR product only in the germline. gDNA = germline testis DNA and sDNA = somatic intestine DNA, isolated from the single male worm from which the genome sequences were derived. The 1,006 bp PCR product present in all lanes represents a control PCR corresponding to a single copy locus (miR-279) present in both the germline and somatic genomes.

D. PCR data confirm the elimination of DNA in 17 additional independent loci in the *A. suum* somatic genome. The PCR strategy illustrated in B and C was applied to these loci. See also Figure S1 and Table S2.

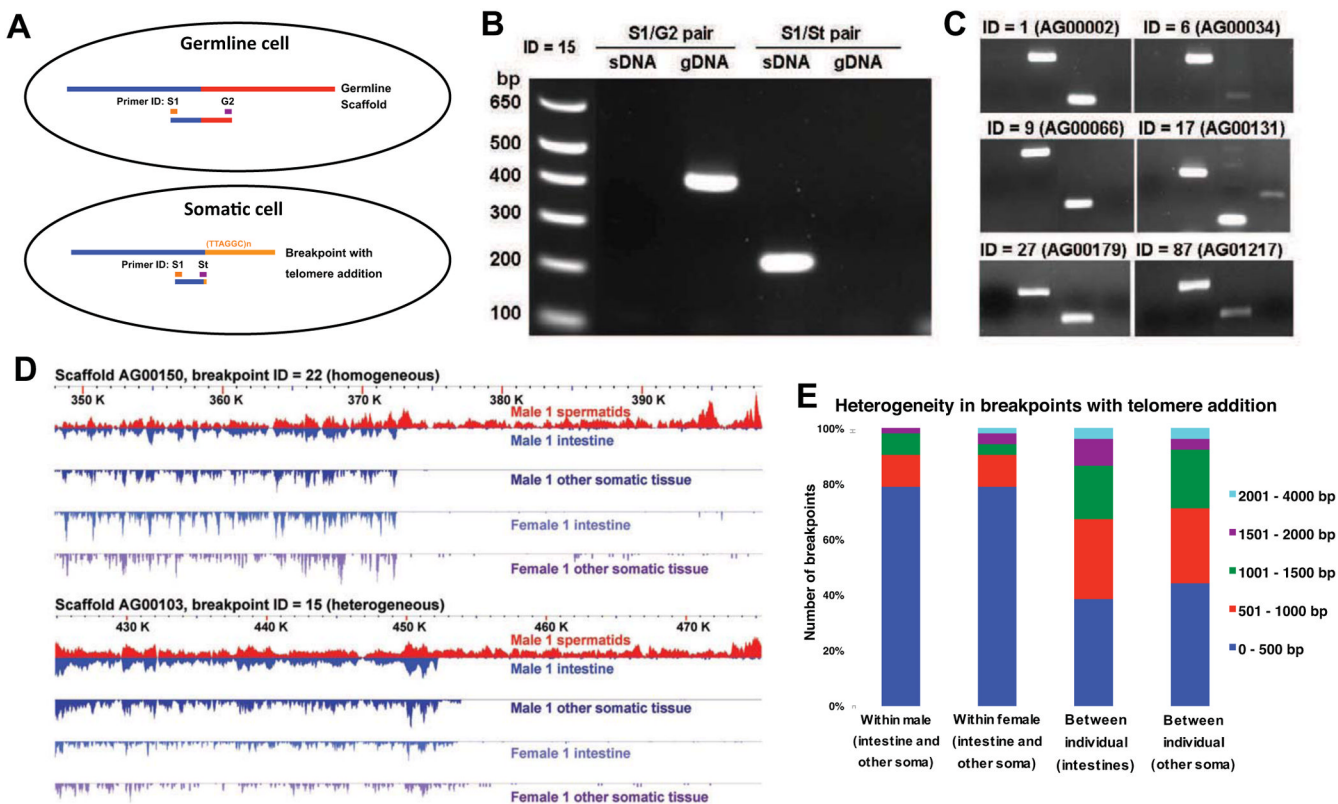


Figure 3. *A. suum* chromosome breaks with telomere addition in somatic cells
A. The PCR strategy used to verify telomere sequence addition in the somatic cells. Primer St (Somatic telomere) is a hybrid primer consisting of 3' nucleotides corresponding to the unique somatic sequence and 5' nucleotides corresponding to telomeric sequence [(TTAGGC)_n] (see Supplementary Experimental Procedures for primers sequences).
B. PCR data confirm telomere addition in the *A. suum* somatic tissues at breakpoint 15. Primers are defined in Fig. 3A and genomic DNA sources defined as in Fig. 2C.
C. PCR data confirm the telomere addition at 6 additional independent loci with chromosome breaks.
D. Heterogeneity in breakpoints with telomere addition. Two breakpoints with telomere addition are illustrated. Note that the exact breakpoint for one of the loci in different somatic tissues varies (intestine and carcass), particularly between individuals.
E. Overall heterogeneity in breakpoints with telomere addition. For these 52 breakpoints, we compared the genomes of pairs of somatic tissues (intestine and other somatic) from the same individual and between individuals and measured the difference in the position of the breakpoints identified.
See also Figure S2 and Table S3.

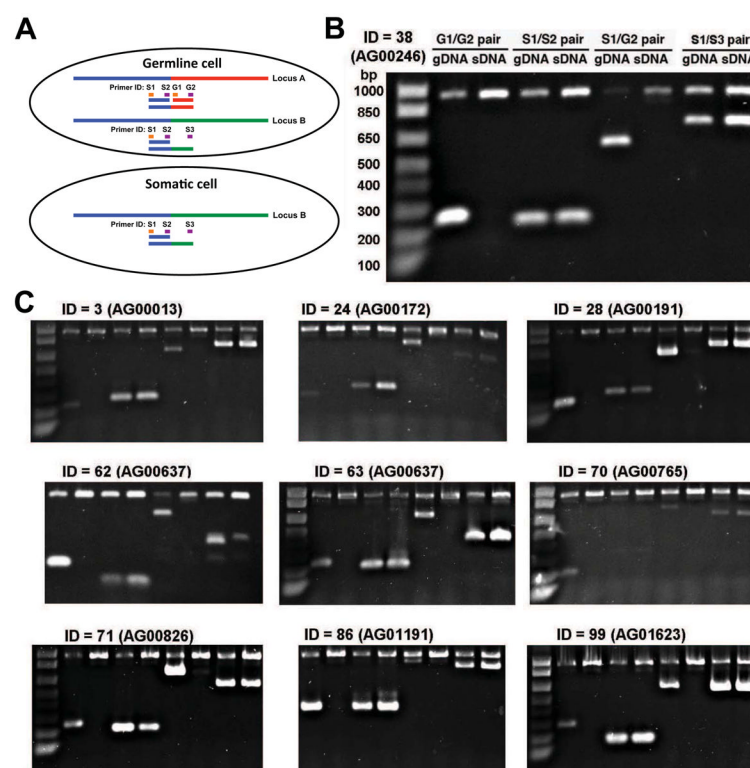


Figure 4. Loss of one member of duplicated, rearranged loci

A. Loss of one of two similar germline loci in the somatic genome. Illustration shows two germline loci in a germline cell containing common sequences (> 97% identical) (blue line), divergent sequence (green or red lines), and the loss of one germline locus in somatic cells following chromatin diminution. Primers and PCR strategy used to verify loci in the germline and somatic genomes are shown.

B. PCR data confirm locus A is present in germline cells but lost in somatic cells. Locus B is present in both germline and somatic cells.

C. Additional PCR data for other loci demonstrating the loss of one member of duplicated, rearranged loci.

See also Table S4.

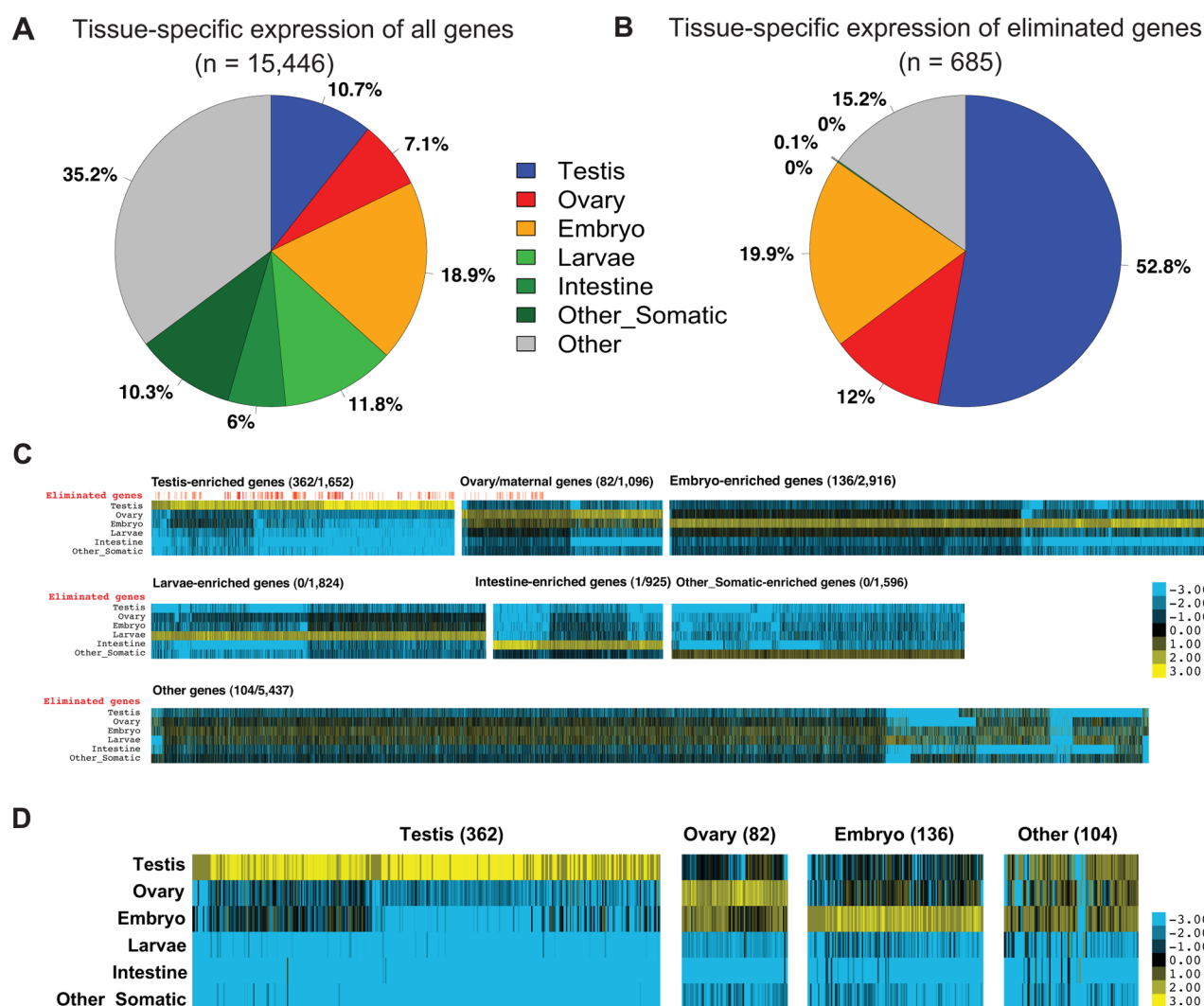


Figure 5. Eliminated *A. suum* genes are primarily expressed in the germline

A. RNA expression of *A. suum* genes (n = 15,446) in different tissues. Gene expression enrichment was categorized by comparing RNA-seq data (Table S1) using reads per kilobase of template per million mapped reads (RPKM).

B. The 685 eliminated *A. suum* genes are highly expressed in the germline and early embryogenesis.

C. Expression Heatmap for all *A. suum* genes. Shown are expression heatmaps for different groups of genes illustrated in Fig. 5A. For each gene, the colors represent log2 values of fold changes to the average expression level (RPKM) for a gene in different stages. For each group of genes, the eliminated/total number of genes is indicated and a red vertical line above the heatmap marks the genes eliminated (see Fig. 5D).

D. The expression profiles of eliminated *A. suum* genes. See Fig. 5C for legend. Note that for the 104 genes in the *Other* group, the majority of them are expressed in testis, ovary, and the early embryo.

See also Figure S3 and Table S5–6.

Table 1*Ascaris suum* germline and somatic genome assemblies

<i>Ascaris</i> genomes	Features	Germline	Somatic	Jex et. al ¹
Assemblies	Estimated genome size (Mb) ²	~ 334	~ 291	~ 309
	Total number of bp assembled (bp)	265,545,801	251,265,282	272,782,664
	N50 of scaffolds (bp); N50 number	290,558; 260	65,087; 1,011	407,899; 179
	N90 of scaffolds (bp); N90 number	48,674; 1,100	11,448; 4,399	80,017; 748
	Number of scaffolds (>= 2,000 bp)	31,538 (2,186)	37,686 (7,692)	29,831 (1,618)
	Maximum length of scaffold (bp)	1,465,500	600,478	3,795,215
	N50 of contigs (bp); N50 number	49,549; 1,510	36,306; 1,925	23,038; 3,512
	N90 of contigs (bp); N90 number	11,178; 5,601	7,566; 7,407	5,913; 11,869
Protein-coding genes	Putative coding gene number	15,446	14,761	18,542
	Average gene size (bp)	9,467	9,473	6,536
	Average coding sequence length (bp)	1,119	1,128	983
	Average exon number per gene	8.4	8.4	6
	Average exon length (bp)	201	201	153
	Average intron length (bp)	1,056	1,050	1,081
Non-coding RNAs	Ribosomal RNAs (rRNAs, copies for 18s-5.8s-26s) ²	~ 500	~ 500	NA
	Splice leader RNAs (including 5s rRNA) ²	~ 265	~ 265	NA
	Transfer RNAs (tRNAs) (+ tRNA pseudogenes)	383 (+ 31)	172 (+ 13)	255 (+ 16)
	microRNAs (miRNAs)	100	100	100
Functional coverage	% of cDNA contigs >= 200 nt present (total 58,085) ³	91.9%	88.2%	92.2%
	% of cDNA bases present (total 58.1 Mb)	97.6%	94.6%	98.0%
	% of unique small RNA reads mapped (total 20.2M) ⁴	81.3%	70.7%	76.3%
	% of all small RNA reads mapped (total 690.7M) ⁴	89.1%	80.5%	78.6%

¹ See Jex et al., 2011.² Genome size and repetitive RNA copy number estimation based on coverage (see Supplementary Experimental Procedures).³ For the de-novo assembled cDNA contigs, > 90% of the sequence mapped back to the genome using BLAT.⁴ See Wang et al., 2011.

See also Figure S4 and Table S1.